Amendments to the Claims:

 (Currently Amended) A method of treating urge, stress or mixed-urinary incontinence comprising administration of an effective amount of a compound of formula IA-IF having the following structure:

wherein:

or C1-C6 haloalkyl:

the carbon atom designated * is in the R or S configuration;

$$\begin{split} R^1 & is \ C_1\text{-}C_6 \ alkyl, \ C_2\text{-}C_6 \ alkenyl, \ C_2\text{-}C_6 \ alkynyl, \ C_3\text{-}C_6 \ cycloalkyl \ or \ C_4\text{-}C_7 \ cycloalkyl \ alkyl, \\ each \ of \ which \ is \ optionally \ substituted \ with 1 \ to 3 \ substitutents \ independently \ selected \ at \\ each \ occurrence \ thereof \ from \ C_1\text{-}C_3 \ alkyl, \ halogen, \ aryl, \ -CN, \ -OR^9 \ and \ -NR^9R^{10} \ ; \\ R^2 \ is \ H, \ C_1\text{-}C_6 \ alkyl, \ C_2\text{-}C_6 \ alkynyl, \ C_3\text{-}C_6 \ cycloalkyl, \ C_4\text{-}C_7 \ cycloalkyl \ alkyl \ and \ cycloalkyl \$$

 R^3 is H, halogen, $-OR^{11}$, $-S(O)R^{12}$, $-S(O)_n$ $NR^{11}R^{12}$, -CN, $-C(O)R^{12}$, $-C(O)NR^{11}R^{12}$, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_6 cycloalkyl, C_4 - C_7 cycloalkylalkyl, -O(phenyl) or -O(benzyl), wherein each of -O(phenyl) and -O(benzyl) is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, or C_1 - C_4 alkoxy, or wherein R^3 is a C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_6 cycloalkyl or C_4 - C_7 cycloalkylalkyl group, then said group is optionally substituted with from 1 to 3 substituents selected independently at each occurrence thereof from C_1 - C_3 alkyl, halogen, aryl, -CN, $-OR^9$ and $-NR^9R^{10}$:

provided that for compounds of formula IA, R³ is C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkenyl, C₃-C₆ cycloalkyl or C₄-C₇ cycloalkylalkyl, each of which is optionally substituted with from 1 to 3 substituents selected independently at each occurrence thereof from C₁-C₃ alkyl, halogen, aryl, -CN, -OR⁹ and -NR⁹R¹⁰;

provided that for compounds of formula IB, R^3 is - O(phenyl), -O(benzyl), -OC(O) R^{13} or - $S(O)_n R^{12}$, each of -O(phenyl) and -O(benzyl) is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, or C_1 - C_4 alkoxy;

 R^4 is H, halogen, -OR 11 , -S(O)_nR 12 , -S(O)NR $^{11}R^{12}$, -CN, -C(O)R 12 , -C(O)NR $^{11}R^{12}$, -NR $^{11}R^{12}$, C1-C6 alkyl, C2-C6 alkenyl, C2-C6 alkynyl, C3-C6 cycloalkyl, C4-C7 cycloalkylalkyl, O(phenyl) or -O(benzyl), wherein each of -O(phenyl) and -O(benzyl) is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C1-C4 alkyl, C1-C4 haloalkyl or C1-C4 alkoxy and wherein R^4 is a C1-C4 alkyl, C2-C6 alkenyl, C2-C6 alkynyl, C3-C6 cycloalkyl or C4-C7 cycloalkylalkyl group, then said group is optionally substituted with from 1 to 3 substituents selected independently at each occurrence thereof from C1-C3 alkyl, halogen, aryl, -CN, - OR 9 and -NR $^9R^{10}$.

provided that for compounds of formula IC, R_4 is C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkenyl, C_3 - C_6 cycloalkyl, or C_4 - C_7 cycloalkylalkyl, each of which is optionally substituted with from 1 to 3 substituents selected independently at each occurrence thereof from C_1 - C_3 alkyl, halogen, aryl, -CN, -OR 9 and -NR 9 R 10 , or R^5 and R^6 or R^6 and R^7 may be -0-C(R^{12})₂-O-; provided that for compounds of formula ID, R^4 is -O(phenyl), -O(benzyl), -OC(O)R 13 , - $NR^{11}R^{12}$ or -S(O)_R R^{12} , each of -O(phenyl) and -O(benzyl) is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C_1 - C_4 alkyl, C_1 - C_4 alloalkyl, or C_1 - C_4 alkoxy;

 R^5,R^6 and R^7 in compounds of each of the formulae IA, IB, IC, ID, IE and IF are each independently H, halogen, $-OR^{11},-S(O)_nR^{12},-CN,-C(O)R^{12},-NR^{11}R^{12},-C(O)NR^{11}R^{12},-NR^{11}C(O)R^{12},-NR^{11}C(O)_2R^{12},-NR^{11}C(O)NR^{12}R^{13},\,C_{1}-C_6$ alkyl, C_2-C_6 alkenyl, C_2-C_6 alkenyl, C_3-C_6 cycloalkyl or C_4-C_7 cycloalkylalkyl, wherein each of R^5,R^6 and R^7 is a C_1-C_6 alkyl, C_2-C_6 alkenyl, C_2-C_6 alkynyl, C_3-C_6 cycloalkyl or C_4-C_7 cycloalkyl group, then

said group is optionally substituted with from 1 to 3 substituents selected independently at each occurrence thereof from C_1 - C_3 alkyl, halogen, aryl, -CN, -OR⁹ and -NR⁹R¹⁰, or R⁵ and R⁶ or R⁶ and R⁷ may be -0-C(R¹²)₂-O-;

provided that for compounds of formula IE at least one of \mathbb{R}^5 or \mathbb{R}^7 is fluoro, chloro, or methyl;

or R^7 and R^6 are each independently -O-C(R^{12})₂-0- in compounds of the formulae IE, but only where R^2 is fluoro, chloro or methyl;

or R^7 and R^6 can independently also be -O-C(R^{12})₂-0- in compounds of the formulae IE, but only where R^7 is fluoro, chloro or methyl;

R⁸ is H, halogen, or OR¹¹, provided that for compounds of formula IF, R⁸ is halogen;
R⁹ and R¹⁰ are each independently H, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxyalkyl, C₃C₆ cycloalkyl, C₄-C₇ cycloalkylalkyl, -C(O)R¹³, phenyl or benzyl, where phenyl or benzyl is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C₁-C₄ alkyl, C₁-C₄ haloalkyl, or C₁-C₄ alkoxy; or R⁹ and R¹⁰ are taken together with the nitrogen to which they are attached to form piperidine, pyrrolidine, piperazine, N-methylpiperazine, morpholine, or thiomorpholine;
R¹¹ is H, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxyalkyl, C₃-C₆ cycloalkyl, C₄-C₇ cycloalkylalkyl, -C(O)R¹³, phenyl or benzyl, where R¹¹ is a C₁-C₄ alkyl, phenyl or benzyl group, then said group is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C₁-C₄ alkyl, C₁-C₄ haloalkyl, or C₁-C₄ alkoxy.

 R^{12} is H, amino, C_1 - C_4 alkyl, $(C_1$ - C_4 alkyl)amino, C_1 - C_4 haloalkyl, C_1 - C_4 alkoxyalkyl, C_3 - C_6 cycloalkyl, C_4 - C_7 cycloalkyl, phenyl or benzyl, where phenyl or benzyl is optionally substituted from 1 to 3 times with a substituent selected independently from halogen, cyano, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl and C_1 - C_6 alkoxy;

or R¹¹ and R¹² are taken together with the nitrogen to which they are attached to form piperidine, pyrrolidine, piperazine, N-methylpiperazine, morpholine, or thiomorpholine; provided that only one of R⁹ and R¹⁰ or R⁹ and R¹⁰ are taken together with the nitrogen to which they are attached to form piperldine, pyrrolidine, piperazine, N-methylpiperazine, morpholine.

- R13 is C1-C4 alkyl, C1-C4 haloalkyl or phenyl;
- n is 0, 1, or 2, and;
- aryl is phenyl which is optionally substituted 1-3 times with halogen, cyano, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl and C_1 - C_4 alkoxy,
- or an oxide thereof, a pharmaceutically acceptable salt thereof, a solvate thereof, or prodrug thereof.
- 2. (Original) A method of claim 1, wherein R¹ is C₁-C₃ alkyl.
- 3. (Original) A method of claim 2, wherein R1 is CH3.
- 4. (Original) A method of claim 1, wherein R² is H, C₁-C₄ alkyl or C₁-C₆ haloalkyl.
- 5. (Original) A method of claim 4, wherein R² is H or CH₃.
- 6. (Original) A method of claim 1, wherein R³ is H or R³ is C₁-C₄ alkyl, C₃-C₆ cycloalkyl or C₄-C₇ cycloalkylalkyl, each of which is optionally substituted with from 1 to 3 substituents selected independently at each occurrence thereof from C₁-C₃ alkyl, halogen, aryl, -CN, -OR9 and NR³R¹0, or R³ is -O(phenyl) or -O(benzyl) optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C₁-C₄ alkyl, C₁-C₄ haloalkyl, or C₁-C₄ alkoxy.
- 7. (Original) A method of claim 6, wherein R³ is methyl, ethyl, propyl, or isopropyl.
- 8. (Original) A method of claim 6, wherein R³ is -O(phenyl) or -O-CH₂-(phenyl), each of which is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C₁-C₄ alkyl, C₁-C₄ haloalkyl, or C₁-C₄ alkoxy.
- 9. (Original) A method of claim 6, wherein R3 is H.

- 10. (Original) A method of claim 1, wherein R⁴ is H, or R⁴ is -NR¹¹R¹² or R⁴ is C₁-C₄ alkyl, C₃-C₆ cycloalkyl or C₄-C₇ cycloalkylalkyl, each of which is optionally substituted, or wherein R⁴ is -O(phenyl) or -O(benzyl), each of which is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C₁-C₄ alkyl, C₁-C₄ haloalkyl, or C₁-C₄ alkoxy.
- 11. (Original) A method of claim 10, wherein R⁴ is methyl, ethyl, propyl, or isopropyl.
- 12. (Original) A method of claim 10, wherein R⁴ is -O(phenyl) or -O(CH₂)phenyl, each of which is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C₁-C₄ alkyl, C₁-C₄ haloalkyl, or C₁-C₄ alkoxy.
- 13. (Original) A method of claim 10, wherein R4 is H.
- 14. (Original) A method of claim 1, wherein R^3 and R^4 are each H or wherein R^3 and R^4 are each halogen.
- 15. (Original) A method of claim 1, wherein one of R3 and R4 is H and the other is CH3.
- 16. (Original) A method of claim 1, wherein R⁵, R⁶ and R⁷ are each H, halogen, -OR¹¹, -NR¹¹R¹², C₁-C₆ alkyl and substituted C₁-C₆ alkyl.
- 17. (Original) A method of claim 16, wherein R5, R6 and R7 are each H.
- 18. (Original) A method of claim 16, wherein one of R⁵ or R⁷ is F, Cl or Me and the other of R⁵ or R⁷ and R⁶ are H, halogen, -OR¹¹, -NR¹¹R¹², or optionally substituted C₁-C₆ alkyl.
- 19. (Original) A method of claim 18, wherein R⁵ is F, Cl or Me; and R⁷ is H.
- 20. (Original) The method of claim 18, wherein R5 is F, Cl or Me, and R6 is H.

- 21. (Original) A method of claim 1, wherein R8 is halogen.
- 22. (Original) A method of claim 21, wherein R⁸ is fluoro.
- 23. (Original) A method of claim 1, wherein:

$$R^1$$
 is C_1 - C_3 alkyl;

 R^2 is H, C_1 - C_4 alkyl or C_1 - C_6 haloalkyl;

 R^3 is $C_1\text{-}C_4$ alkyl, $C_3\text{-}C_6$ cycloalkyl or $C_4\text{-}C_7$ cycloalkylalkyl, each of which is optionally substituted, or R^3 is -O(phenyl) or -O(benzyl), each of which is optionally substituted, or R^3 is H; R^4 is H, $C_1\text{-}C_4$ alkyl, $C_3\text{-}C_6$ cycloalkyl or $C_4\text{-}C_7$ cycloalkylalkyl, each of which is optionally substituted with from 1 to 3 substituents selected independently at each occurrence thereof from $C_1\text{-}C_3$ alkyl, halogen, aryl, -CN, -OR 9 and -NR $^9R^{10}$, or R^4 is -NR $^{11}R^{12}$, - O(phenyl) or -O(benzyl), wherein said -O(phenyl) or -O(benzyl), is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, $C_1\text{-}C_4$ alkyl, $C_1\text{-}C_4$ haloalkyl, or $C_1\text{-}C_4$ alkoxy;

or R3 and R4 are each halogen;

 R^5 , R^6 and R^7 are each H, halogen, $-OR^{11}$, $-NR^{11}R^{12}$, optionally substituted C_1 - C_6 alkyl, or one of R^5 and R^7 is C_1 , C_1 or Me and the other of R^5 and R^7 and R^6 is H, halogen, $-OR^{11}$, $-NR^{11}R^{12}$, C_1 - C_6 alkyl or substituted C_1 - C_6 alkyl.

24. (Original) A method of claim 23, wherein:

R1 is CH3;

R2 is H or CH3;

 R^3 is H, F, methyl, ethyl, propyl, isopropyl, -O(phenyl) or -0-CH₂-(phenyl), wherein said -O(phenyl) or -0-CH₂-(phenyl) is optionally substituted from 1 to 3 times with a substituent selected independently at each occurrence thereof from halogen, cyano, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, or C_1 - C_4 alkoxy;

 R^4 is H, F methyl, ethyl, propyl, isopropyl, -O(phenyl) or -0-CH₂-(phenyl), wherein said -O(phenyl) or -0-CH₂-(phenyl) is optionally substituted from 1 to 3 times with a substituent

selected independently at each occurrence thereof from halogen, cyano, C₁-C₄ alkyl, C₁-C₄ haloalkyl, or C₁-C₄ alkoxy;

 R^5 , R^6 and R^7 are each H or R^5 is F, CI or Me, or one of R^6 or R^7 is H and the other of R^6 and R^7 is halogen, $-OR^{11}$, $-NR^{11}R^{12}$, or optionally substituted C_1 - C_6 alkyl.

- 25. (Original) A method of claim 23, wherein R8 is halogen.
- 26. (Original) A method according to claim 1, wherein the carbon atom designated * is in the R configuration.
- 27. (Original) A method according to claim 1, wherein the carbon atom designated * is in the S configuration.
- 28. (Original) A method comprising a mixture of stereoisomeric compounds of claim 1 wherein the carbon atom designated * is in the S or R configuration.
- 29. (Currently Amended) A method according to claim 1, wherein the compound is selected from the group consisting of:
 - 2,7-dimethyl-4-phenyl-1,2,3,4-tetrahydroisoquinoline;
 - 4-(4-methoxy)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoquinoline;
 - 2,7-dimethyl-4-(4-fluoro)phenyl-1,2,3,4-tetrahydroisoquinoline;
 - 2,7-dimethyl-4-(3-fluoro)phenyl-1,2,3,4-tetrahydroisoquinoline;
 - 4-(3,4-difluoro)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoquinoline;
 - 2,7-dimethyl-4-(4-fluoro-3-methyl)phenyl-1,2,3,4-tetrahydroisoquinoline;
 - 4-(3-chloro-4-fluoro)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoquinoline;
 - 4-(3-chloro)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoguinoline;
 - 2,7-dimethyl-4-(4-methyl)phenyl-1,2,3,4-tetrahydroisoquinoline;
 - 2.7-dimethyl-4-(3-fluoro-4-methyl)phenyl-1,2,3,4-tetrahydroisoguinoline;
 - 4-(4-chloro)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoquinoline;
 - 4-(4-chloro-3-fluoro)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoquinoline;

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4-(3,4-dichloro)phenyl-2,7-dimethyl-1,2,3,4-tetrahydroisoquinoline;
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7-ethyl-2-methyl-4-phenyl-1,2,3,4-tetrahydroisoquinoline;

4-(3.4-difluoro)phenyl-7-ethyl-2-methyl-1,2,3,4-tetrahydroisoguinoline;

7-fluoro-4-(4- methoxy)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;

7-fluoro-4-(3-fluoro-4-methoxy)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;

7- fluoro-4-(3-fluoro-4-methyl)phenyl-2-methyl-1,2,3,4- tetrahydroisoquinoline;

7-fluoro-4-(4-chloro-3-fluoro)phenyl-2-methyl-1,2,3,4- tetrahydroisoquinoline;

4-(3,4-difluoro)phenyl-7-fluoro-2-methyl-1,2,3,4-tetrahydroisoquinoline;

4-(3-chloro)phenyl-7-fluoro-2-methyl-1,2,3,4-tetrahydroisoquinoline;

7-cvano-2-methyl-4-phenyl-1,2,3,4-tetrahydroisoguinoline;

2-methyl-4-phenyl-7-trifluoromethyl-1,2,3,4-tetrahydroisoquinoline;

4-phenyl-1,2,7-trimethyl-1,2,3,4-tetrahydroisoquinoline:

4-(4-chloro)phenyl-1,2-dimethyl-1,2,3,4-tetrahydroisoquinoline;

4-(3,4-difluoro)phenyl-1,2-dimethyl-1,2,3,4-tetrahydroisoquinoline;

4-phenyl-2.7.8-trifluoromethyl-1,2,3,4-tetrahydrolsoguinoline;

2.7-dimethyl-8-fluoro-4-phenyl-1.2.3.4-tetrahydroisoguinoline;

2.8-dimethyl-7-fluoro-4-phenyl-1.2.3.4-tetrahydroisoguinoline;

2,7-dimethyl-8-methoxy-4-phenyl-1,2,3,4-tetrahydroisoguinoline;

2,7-dimethyl-8-hydroxy-4-phenyl-1,2,3,4-tetrahydroisoquinoline;

2-methyl-4-phenyl-7-trifluoromethoxy-1,2,3,4-tetrahydroisoquinoline;

4-(3,4-difluoro)phenyl-7-methoxy-2-methyl-1,2,3,4-tetrahydroisoquinoline;

4-(4-fluoro-3-methyl)phenyl-7-methoxy-2-methyl-1,2,3,4-tetrahydroisoguinoline;

4-(3-fluoro-4-methyl)phenyl-7-methoxy-2-methyl-1,2,3,4-tetrahydroisoquinoline;

7-methoxy-4-(3-methyl)phenyl-2-methyl-1,2,3,4-tetrahydroisoguinoline;

2-methyl-7-phenoxy-4-phenyl-1,2,3,4-tetrahydroisoguinoline;

7-(4-methoxy)phenoxy-2-methyl-4-phenyl-1,2,3,4-tetrahydroisoguinoline;

7-benzyloxy-2-methyl-4-phenyl-1,2,3,4-tetrahydroisoguinoline;

7-hvdroxy-2-methyl-4-(3-methyl)phenyl-1,2,3,4-tetrahydroisoguinoline;

4-(3-fluoro-4-methyl)phenyl-7-hydroxy-2-methyl-1,2,3,4-tetrahydroisoguinoline;

4-(4-fluoro-3-methyl)phenyl-7-hydroxy-2-methyl-1,2,3,4-tetrahydrolisoguinoline;

- 4-(3,4-difluoro)phenyl-7-hydroxy-2-methyl-1,2,3,4-tetrahydroisoguinoline;
- 4-(3-cyano)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;
- 2.8-dimethyl-4-phenyl-1,2,3,4-tetrahydroisoguinoline;
- 2.8-dimethyl-4-(4-fluoro)phenyl-1,2,3,4-tetrahydroisoguinoline;
- 4-(3.4-difluoro)phenyl-2.8-dimethyl-1.2.3.4-tetrahydroisoguinoline;
- 4-(3.5-difluoro)phenyl-2.8-dimethyl-1,2,3,4-tetrahydroisoguinoline;
- 2.8-dimethyl-4-(3-fluoro)phenyl-1,2,3,4-tetrahydroisoquinoline;
- 2.8-dimethyl-4-(4-fluoro-3-methyl)phenyl-1,2,3,4-tetrahydroisoquinoline;
- 4-(3-chloro-4-fluoro)phenyl-2,8-dimethyl-1,2,3,4-tetrahydrolsoguinoline;
- 4-(3.4-dichloro)phenyl-2.8-dimethyl-1,2,3,4-tetrahydroisoguinoline;
- 4-(3-chloro)phenyl-2.8-dimethyl-1.2.3,4-tetrahydroisoguinoline;
- 4-(4-chloro)phenyl-2.8-dimethyl-1.2.3.4-tetrahydroisoguinoline;
- 4-(4-chloro-3-fluoro)phenyl-2,8-dimethyl-1,2,3,4-tetrahydroisoquinoline;
- 2,8- dimethyl-4-(4-methoxy)phenyl-1,2,3,4-tetrahydroisoguinoline;
- 4-(4-cyano)phenyl-2.8-dimethyl-1,2,3,4-tetrahydroisoguinoline;
- 2.8-dimethyl-4-(4-trifluoromethyl)phenyl-1,2,3,4-tetrahydroisoguinoline;
- 2.8-dimethyl-4-(4-methyl)phenyl-1.2.3.4-tetrahydroisoguinoline;
- 2-methyl- 8-(N-methylamino)methyl-4-phenyl-1,2,3,4-tetrahydroisoquinoline;
- 8-(hydroxy)methyl-2-methyl-4-phenyl-1,2,3,4-tetrahydroisoquinoline;
- 2-methyl-4-phenyl-8-sulfonamide-1,2,3,4-tetrahydroisoguinoline;
- 2-methyl-8-(N-methyl)sulfonamide-4-phenyl-1,2,3,4-tetrahydroisoguinoline;
- 8-methoxy-2-methyl-4-(4-methyl)phenyl-1,2,3,4-tetrahydroisoguinoline;
- 4-(3.5-difluoro)phenyl-8-methoxy-2-methyl-1.2.3.4-tetrahydroisoguinoline;
- 4-(3-chloro)phenyl-8-methoxy-2-methyl-1,2,3,4-tetrahydroisoguinoline;
- 4-(3,4-dichloro)phenyl-8-methoxy-2-methyl-1,2,3,4-tetrahydroisoquinoline;
- 4-(4-chloro-3-fluoro)phenyl-8-methoxy-2-methyl-1,2,3,4-tetrahydroisoquinoline;
- 4-(3-chloro-4-fluoro)phenyl-8-methoxy-2-methyl-1, 2,3,4-tetrahydroisoquinoline;
- 4-(3,5-difluoro)phenyl-2-methyl-1,2,3,4- tetrahydroisoguinoline;
- 4-(3-chloro-5-fluoro)phenyl-2-methyl-1,2,3,4-tetrahydroisoguinoline;
- $\hbox{$4$-(3,5$-difluoro) phenyl-2,7$-dimethyl-1,2,3,4$-tetrahydroisoquino line;}\\$

- 4-(3-chloro-5-fluoro)phenyl-2,7-dimethyl-1,2,3,4-tetrahydrolsoquinoline;
- 2-methyl-4-(3,4,5-trifluoro)phenyl-1,2,3,4-tetrahydroisoquinoline;
- 4-(3- fluoro)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoiine;
- 4-(3-fluoro-4-methyl)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;
- 4-(4-fluoro-3-methyl)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;
- 4-(3.4-difluoro)phenyl-2-methyl-1,2,3,4-tetrahydroisoguinoline;
- 4-(3-chloro)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;
- 4-(4-chloro-3-fluoro)phenyl-2-methyl-1,2,3,4-tetrahydroisoquinoline;
- 4-(3-chloro-4- fluoro)phenyl-2-methyl-1,2,3,4-tetrahydroisoguinoline;
- 4-(3-cyano)phenyl-2-methyl-1,2,3,4-tetrahydroisoguinoline;
- 4-(4-acetanilide)-2-methyl-1,2,3,4-tetrahydroisoguinoline;
- 4-(4-chloro)phenyl-4-fluoro-2-methyl-1,2,3,4-tetrahydroisoquinoline;
- (3,5-difluoro)-4-phenyl-1,2,7-trimethyl-1,2,3,4-tetrahydroisoquinoline;
- (8-fluoro-2-methyl-4-phenyl-1,2,3,4-tetrahydro-7-isoquinolinyl)-N-methylmethanamine;
- (2-methyl-4-phenyl-7-isoquinolinyl)-N-methylmethanamine;
- N-methyl-(2-methyl-4-phenyl-7-isoquinolinyl)-N-methylmethanamine;
- 8-hydroxy-2-methyl-4-phenyl-1,2,3,4-tetrahydro-7-isoquinolinecarbonitrile;
- (2-methyl-4-phenyl-1,2,3,4-tetrahydro-7-isoquinolinyl)methanol; and
- 2-ethyl-4-phenyl-1,2,3,4-tetrahydroisoquinoline; and
- an oxide thereof, a pharmaceutically acceptable salt thereof, a solvate thereof, or prodrug thereof.
- 30. (New) A method of claim 1, wherein the urinary incontinence is urge, stress, or mixed urinary incontinence.